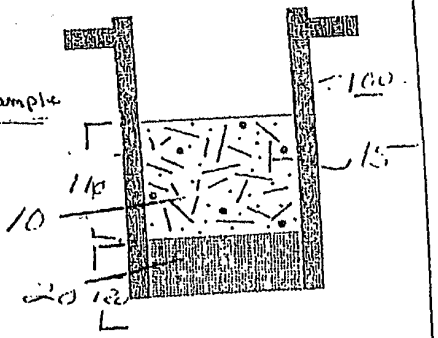
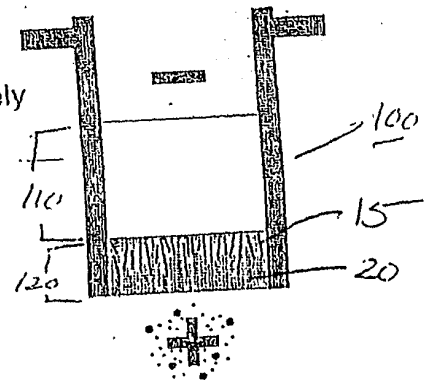
Step 1

- Load DNA analyte sample into microtiter wells



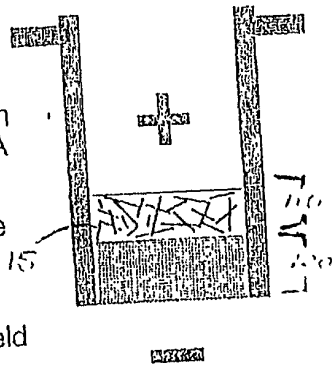
Step 2

- Apply electric field to electrophorese all negatively charged molecules
- The DNA analyte will be captured



Step 3

- Replace buffer
- Reduce sample volume in well if concentration of DNA analyte is also desired
- Apply current to denature hybrid and release DNA analyte from capture oligo
- Apply reversed electric field to electrophoretically elute DNA analyte into the sample volume in the well



Step 4

- Microtiter plate bearing purified and (optionally) concentrated DNA analyte samples ready for further analysis

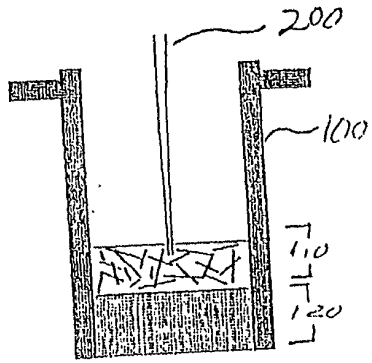


Fig. 1

Capture Probes and Uses Therefor
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Serial No.: Not yet assigned
Atty Docket No.: EXT-070C1
Atty/Agent: Patrick R.H. Waller
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Sample Prep for Sequencing

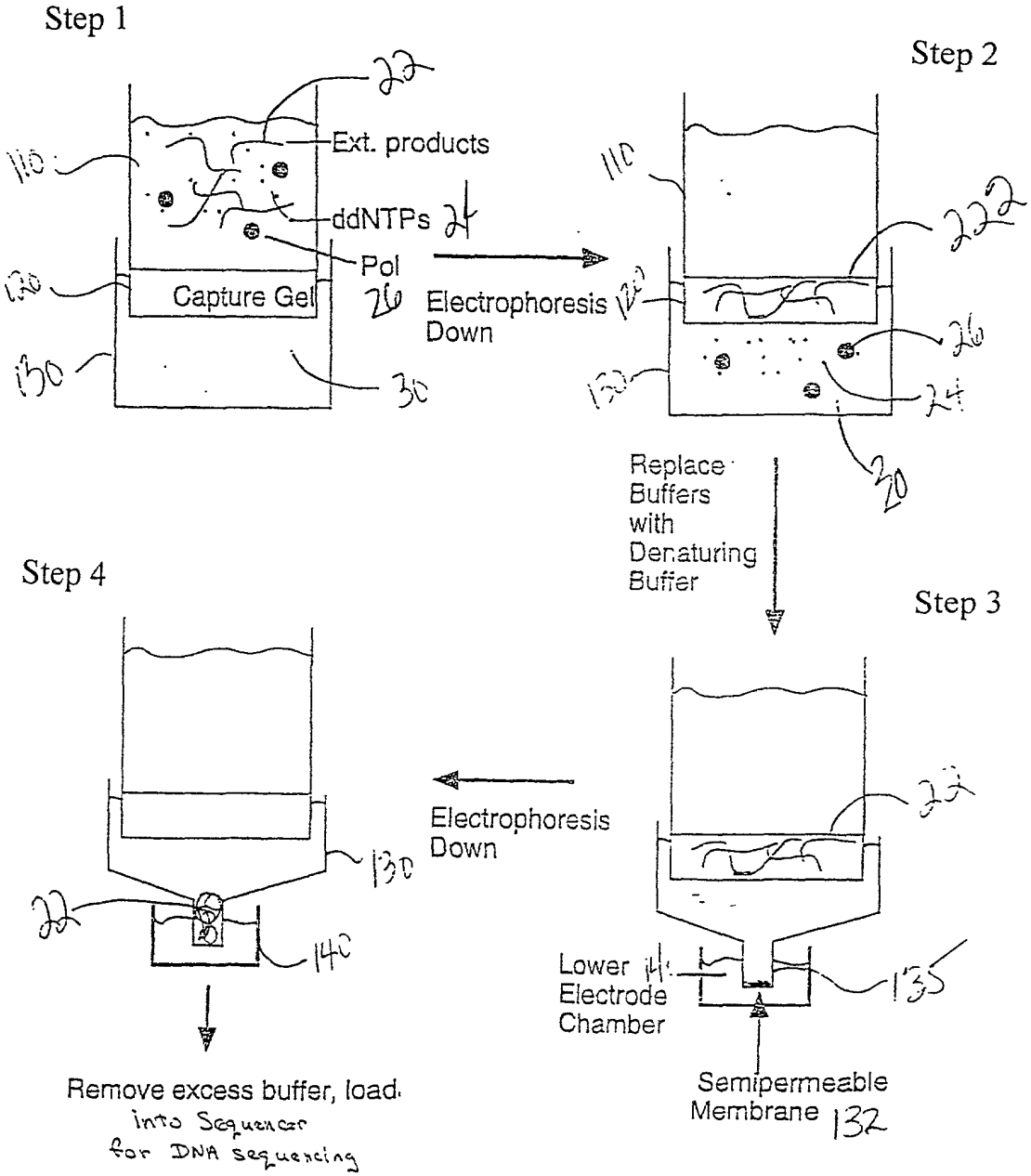

Fig. 2

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Multiplexing: Use of Hybrigel to Purify Products of Multiple Reactions

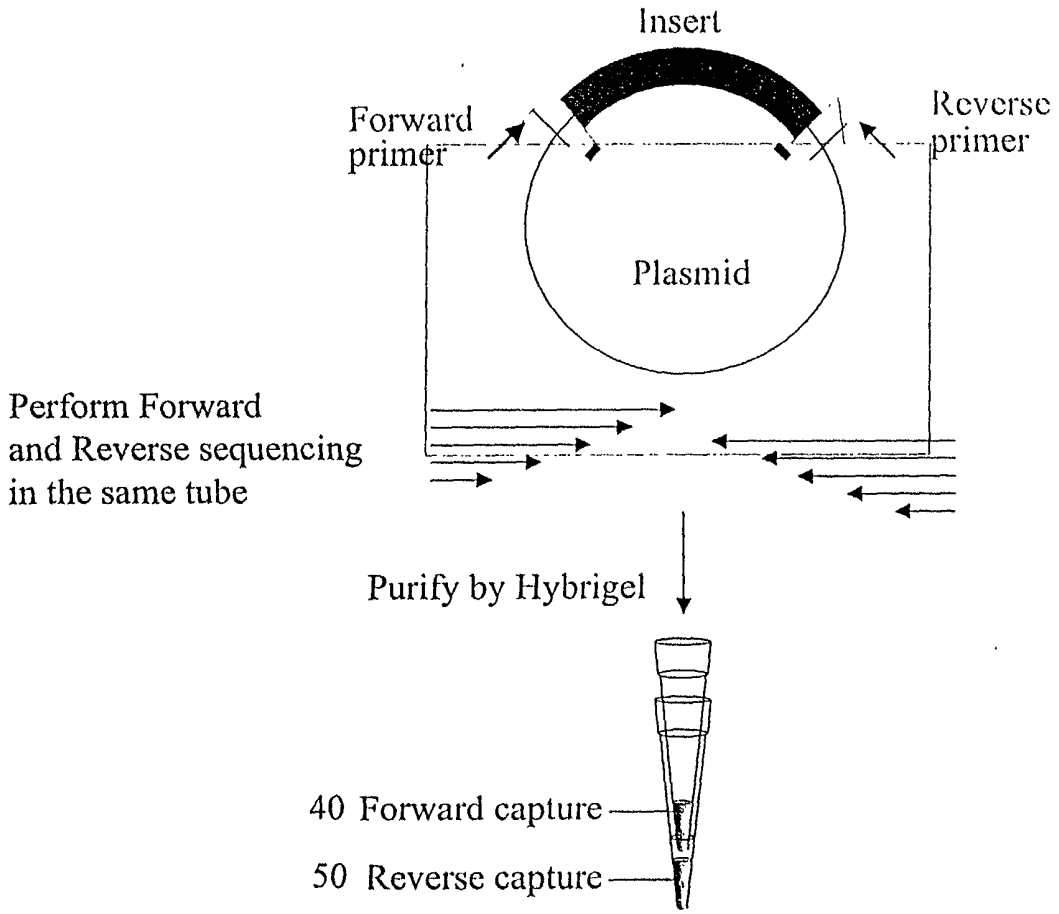

Fig. 3A

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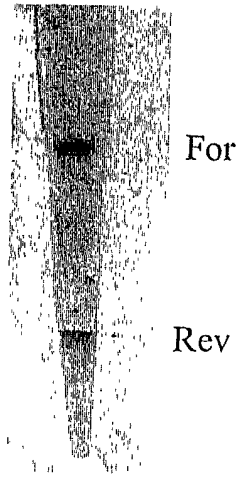

Fig. 3B

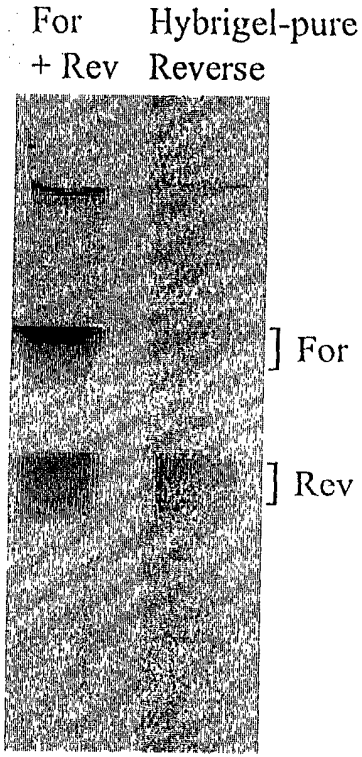

Fig. 3C

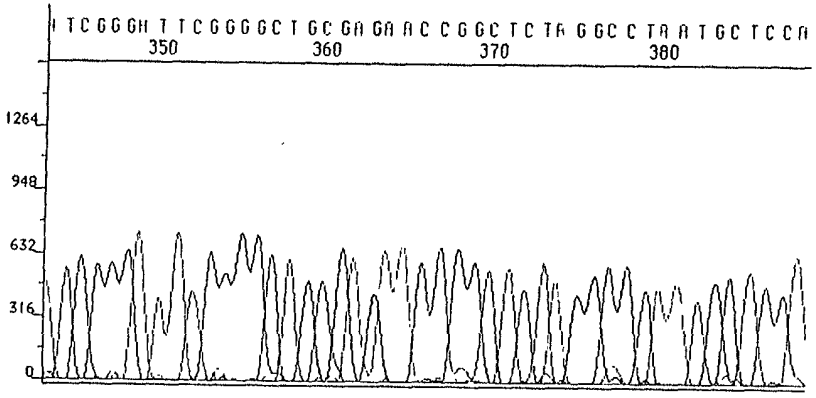

Fig. 3D

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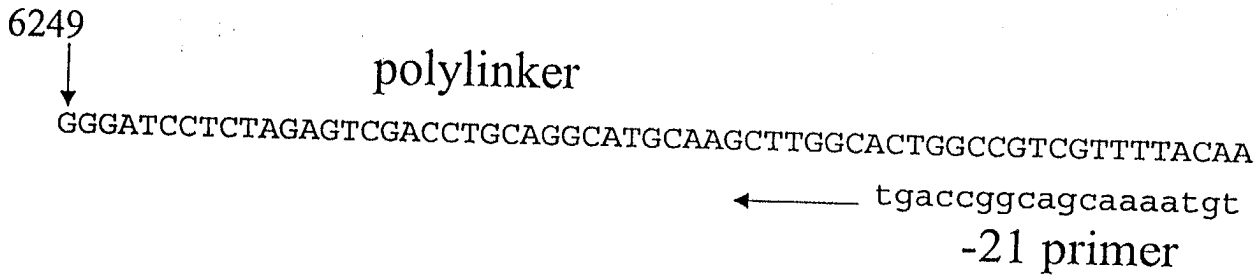

5' ac-GGGATCCTCTAGAGTCG 6249-17ac

5' ac-TGCAGGCATGCAAGCTT 6269-17ac

5' ac-ATGCAAGCTTGGCACTGGCCG 6277-21ac

5' ac-ACTGGCCGTCGTTTTACA 6290-18ac

Fig. 3E

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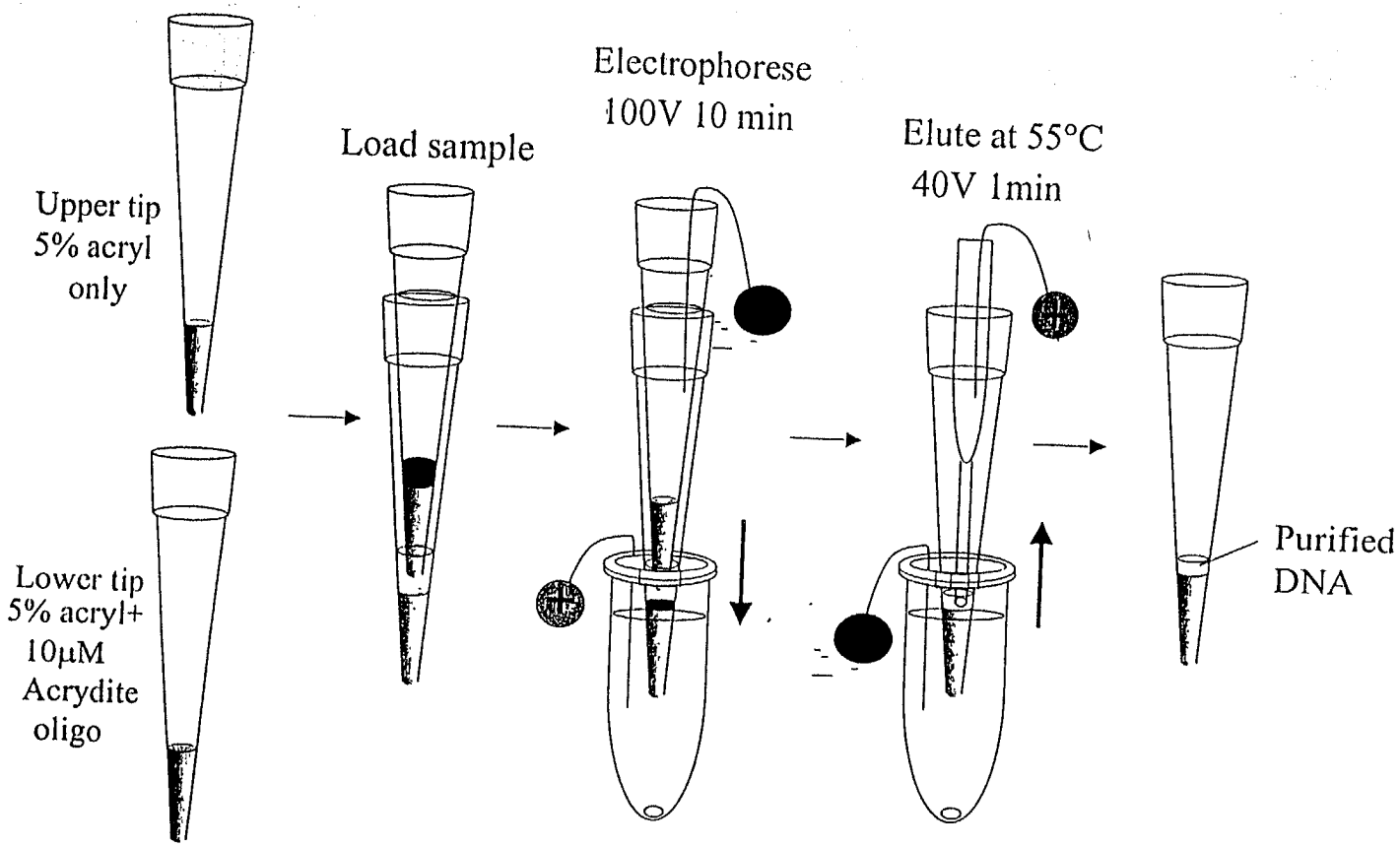

Fig. 4A

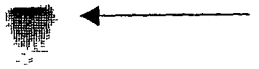

Fig. 4B

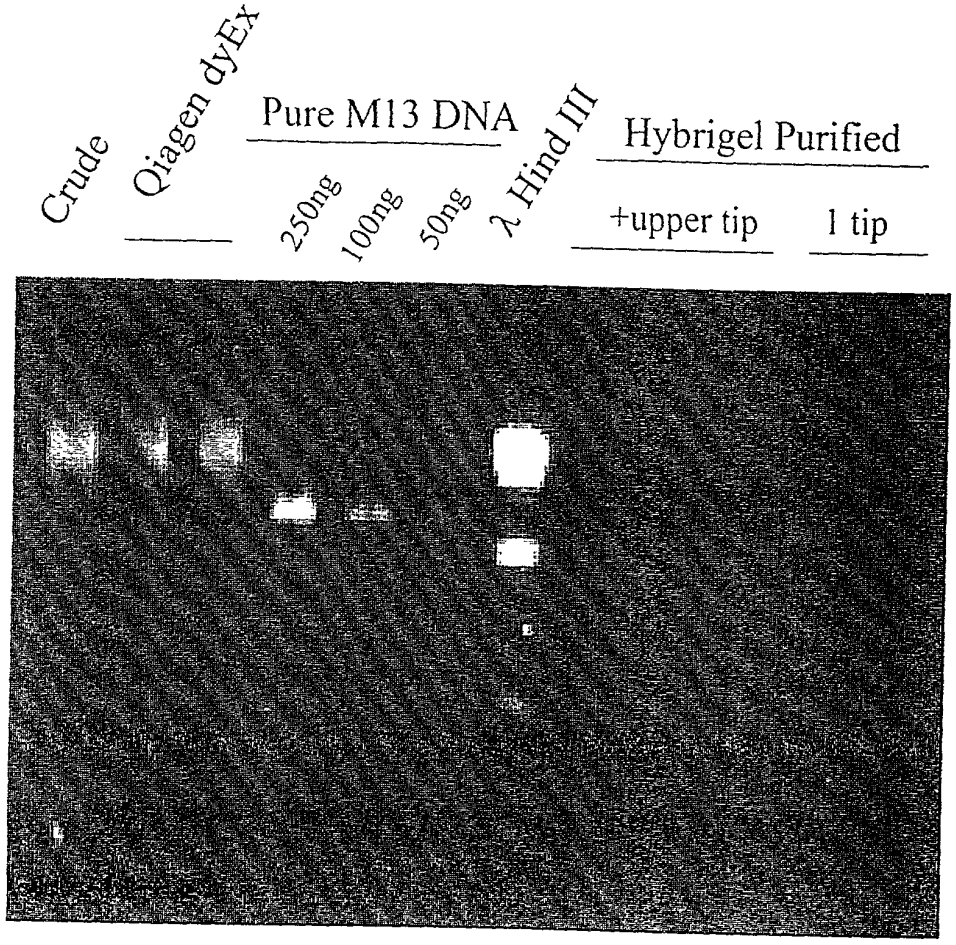
Crude
Qiagen dyEx
Pure M13 DNA
250ng
100ng
50ng
λ Hind III
Hybrigel Purified
+upper tip
1 tip

Fig. 5

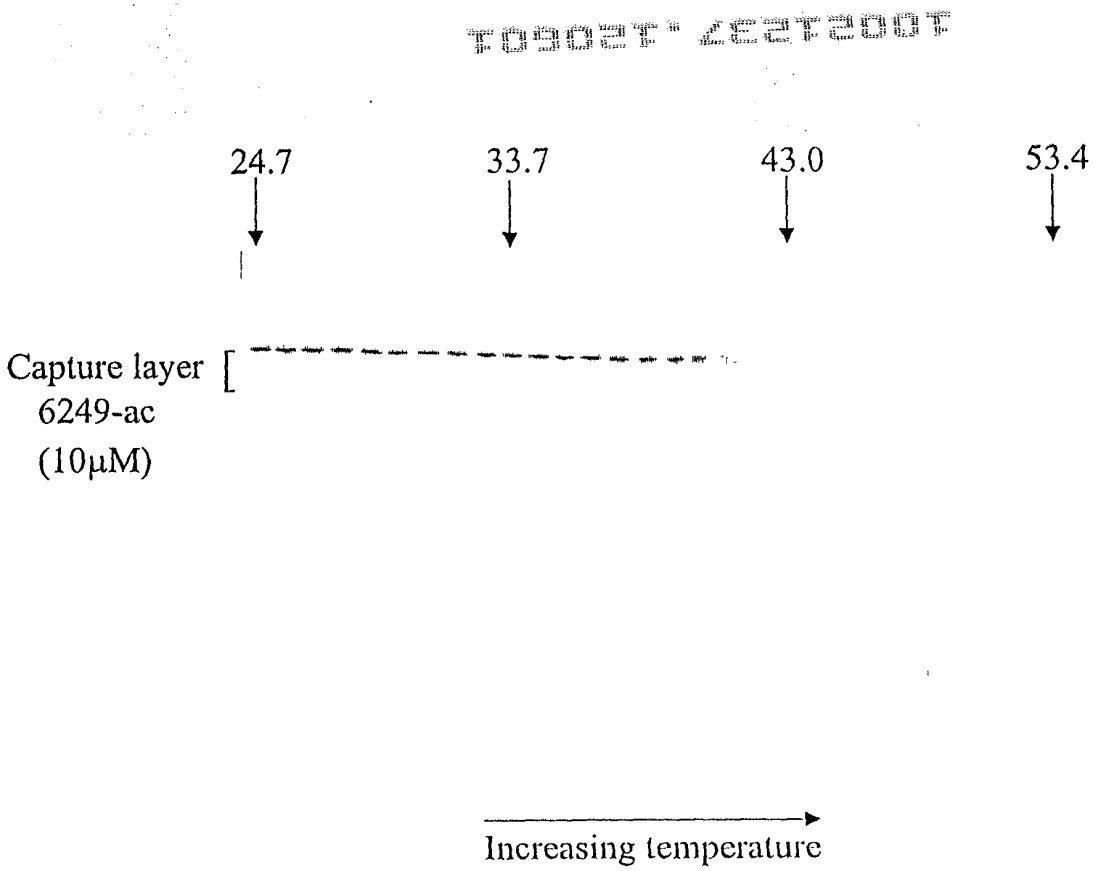
24.7
33.7
43.0
53.4
Capture layer
6249-ac
(10μM)
Increasing temperature

Fig. 6

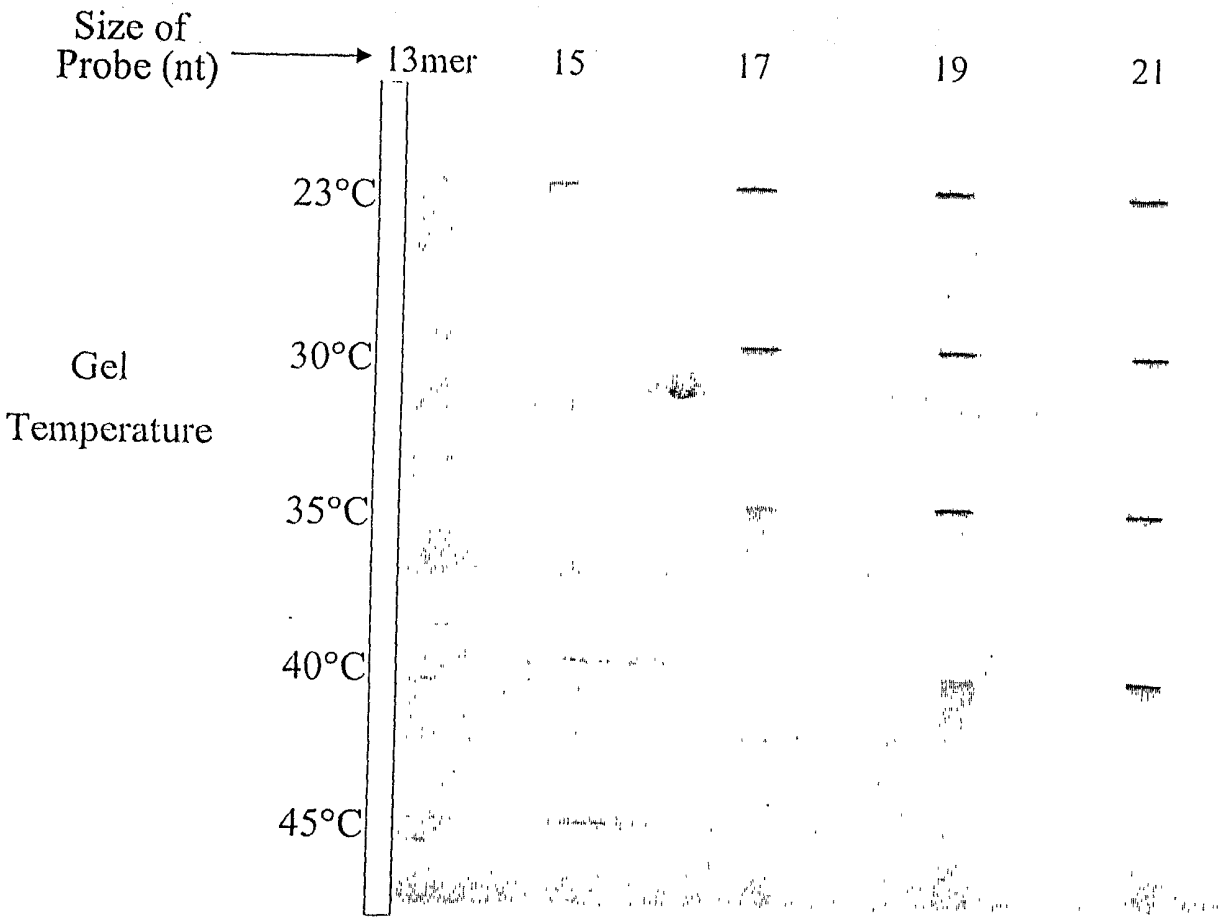

Fig. 7

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